John S K Kauwe

List of Publications by Year in descending order

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38660 24,787 136 50 citations h-index papers

g-index 156 156 156 23824 docs citations times ranked citing authors all docs

11581

135

#	Article	IF	CITATIONS
1	Analysis of highâ€risk pedigrees identifies 11 candidate variants for Alzheimer's disease. Alzheimer's and Dementia, 2022, 18, 307-317.	0.4	10
2	Association between WWOX/MAF variants and dementia-related neuropathologic endophenotypes. Neurobiology of Aging, 2022, 111, 95-106.	1.5	6
3	Knowledge Gaps, Challenges, and Opportunities in Health and Prevention Research for Asian Americans, Native Hawaiians, and Pacific Islanders: A Report From the 2021 National Institutes of Health Workshop. Annals of Internal Medicine, 2022, 175, 574-589.	2.0	40
4	Genome-wide association study of brain arteriolosclerosis. Journal of Cerebral Blood Flow and Metabolism, 2022, 42, 1437-1450.	2.4	2
5	New insights into the genetic etiology of Alzheimer's disease and related dementias. Nature Genetics, 2022, 54, 412-436.	9.4	700
6	The Ramp Atlas: facilitating tissue and cell-specific ramp sequence analyses through an intuitive web interface. NAR Genomics and Bioinformatics, 2022, 4, .	1.5	3
7	Novel Alzheimer Disease Risk Loci and Pathways in African American Individuals Using the African Genome Resources Panel. JAMA Neurology, 2021, 78, 102.	4.5	144
8	GenoRisk: A polygenic risk score for Alzheimer's disease. Alzheimer's and Dementia: Translational Research and Clinical Interventions, 2021, 7, e12211.	1.8	7
9	A comprehensive analysis of the phylogenetic signal in ramp sequences in 211 vertebrates. Scientific Reports, 2021, 11, 622.	1.6	2
10	Alzheimer's disease alters oligodendrocytic glycolytic and ketolytic gene expression. Alzheimer's and Dementia, 2021, 17, 1474-1486.	0.4	37
11	Phylogeography of two marine predators, giant trevally (<i>Caranx ignobilis</i>) and bluefin trevally (<i>Caranx melampygus</i>), across the Indo-Pacific. Bulletin of Marine Science, 2021, 97, 257-280.	0.4	6
12	Alzheimer's Disease Alters Oligodendrocytic Glycolytic and Ketolytic Gene Expression. FASEB Journal, 2021, 35, .	0.2	1
13	<i>De novo</i> genome assembly of the marine teleost, bluefin trevally (<i>Caranx melampygus</i>). G3: Genes, Genomes, Genetics, 2021, 11, .	0.8	2
14	Analysis of genes (TMEM106B, GRN, ABCC9, KCNMB2, and APOE) implicated in risk for LATE-NC and hippocampal sclerosis provides pathogenetic insights: a retrospective genetic association study. Acta Neuropathologica Communications, 2021, 9, 152.	2.4	26
15	Pairwise Correlation Analysis of the Alzheimer's Disease Neuroimaging Initiative (ADNI) Dataset Reveals Significant Feature Correlation. Genes, 2021, 12, 1661.	1.0	5
16	Interaction Between Physical Activity and Genes Related to Neurotrophin Signaling in Late-Life Cognitive Performance: The Cache County Study. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2020, 75, 1633-1642.	1.7	7
17	CUBAP: an interactive web portal for analyzing codon usage biases across populations. Nucleic Acids Research, 2020, 48, 11030-11039.	6.5	7
18	Lingering Taxonomic Challenges Hinder Conservation and Management of Global Bonefishes. Fisheries, 2020, 45, 347-358.	0.6	15

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19	Distinct clinicopathologic clusters of persons with TDP-43 proteinopathy. Acta Neuropathologica, 2020, 140, 659-674.	3.9	29
20	Atypical chemokine receptor ACKR2-V41A has decreased CCL2 binding, scavenging, and activation, supporting sustained inflammation and increased Alzheimer's disease risk. Scientific Reports, 2020, 10, 8019.	1.6	7
21	Codon Pairs are Phylogenetically Conserved: A comprehensive analysis of codon pairing conservation across the Tree of Life. PLoS ONE, 2020, 15, e0232260.	1.1	8
22	Genomeâ€wide association study of rate of cognitive decline in Alzheimer's disease patients identifies novel genes and pathways. Alzheimer's and Dementia, 2020, 16, 1134-1145.	0.4	28
23	Identification and genomic analysis of pedigrees with exceptional longevity identifies candidate rare variants. Neurobiology of Disease, 2020, 143, 104972.	2.1	7
24	Predicting Clinical Dementia Rating Using Blood RNA Levels. Genes, 2020, 11, 706.	1.0	10
25	Failure to detect synergy between variants in transferrin and hemochromatosis and Alzheimer's disease in large cohort. Neurobiology of Aging, 2020, 89, 142.e9-142.e12.	1.5	9
26	RAB10: an Alzheimer's disease resilience locus and potential drug target. Clinical Interventions in Aging, 2019, Volume 14, 73-79.	1.3	37
27	Systematic analysis of dark and camouflaged genes reveals disease-relevant genes hiding in plain sight. Genome Biology, 2019, 20, 97.	3.8	122
28	Association study of rs3846662 with Alzheimer's disease in a population-based cohort: the Cache County Study. Neurobiology of Aging, 2019, 84, 242.e1-242.e6.	1.5	5
29	Relative risk for Alzheimer disease based on complete family history. Neurology, 2019, 92, e1745-e1753.	1.5	45
30	Association of Rare Coding Mutations With Alzheimer Disease and Other Dementias Among Adults of European Ancestry. JAMA Network Open, 2019, 2, e191350.	2.8	58
31	Genetic meta-analysis of diagnosed Alzheimer's disease identifies new risk loci and implicates Aβ, tau, immunity and lipid processing. Nature Genetics, 2019, 51, 414-430.	9.4	1,962
32	Assembly of 809 whole mitochondrial genomes with clinical, imaging, and fluid biomarker phenotyping. Alzheimer's and Dementia, 2018, 14, 514-519.	0.4	14
33	Mitochondria and Alzheimer's Disease: the Role of Mitochondrial Genetic Variation. Current Genetic Medicine Reports, 2018, 6, 1-10.	1.9	45
34	Common DNA Variants Accurately Rank an Individual of Extreme Height. International Journal of Genomics, 2018, 2018, 1-7.	0.8	5
35	Genome-wide association study for variants that modulate relationships between cerebrospinal fluid amyloid-beta 42, tau, and p-tau levels. Alzheimer's Research and Therapy, 2018, 10, 86.	3.0	18
36	Analysis of shared heritability in common disorders of the brain. Science, 2018, 360, .	6.0	1,085

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37	Population genealogy resource shows evidence of familial clustering for Alzheimer disease. Neurology: Genetics, 2018, 4, e249.	0.9	6
38	A Common Variant of IL-6R is Associated with Elevated IL-6 Pathway Activity in Alzheimer's Disease Brains. Journal of Alzheimer's Disease, 2017, 56, 1037-1054.	1.2	44
39	Systems biology approach to late-onset Alzheimer's disease genome-wide association study identifies novel candidate genes validated using brain expression data and Caenorhabditis elegans experiments. , 2017, 13, 1133-1142.		40
40	Genome-wide association study identifies four novel loci associated with Alzheimer's endophenotypes and disease modifiers. Acta Neuropathologica, 2017, 133, 839-856.	3.9	199
41	Transethnic genomeâ€wide scan identifies novel Alzheimer's disease loci. Alzheimer's and Dementia, 2017, 13, 727-738.	0.4	166
42	CSF protein changes associated with hippocampal sclerosis risk gene variants highlight impact of GRN/PGRN. Experimental Gerontology, 2017, 90, 83-89.	1.2	7
43	Sex Differences in Risk for Alzheimer's Disease Related to Neurotrophin Gene Polymorphisms: The Cache County Memory Study. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2017, 72, 1607-1613.	1.7	15
44	A common haplotype lowers PU.1 expression in myeloid cells and delays onset of Alzheimer's disease. Nature Neuroscience, 2017, 20, 1052-1061.	7.1	330
45	Using the Health Belief Model to evaluate Samoan caregiver perceptions for rheumatic heart disease follow-up care. International Journal of Health Promotion and Education, 2017, 55, 148-157.	0.4	2
46	Genome-wide, high-content siRNA screening identifies the Alzheimer's genetic risk factor FERMT2 as a major modulator of APP metabolism. Acta Neuropathologica, 2017, 133, 955-966.	3.9	60
47	Rare coding variants in PLCG2, ABI3, and TREM2 implicate microglial-mediated innate immunity in Alzheimer's disease. Nature Genetics, 2017, 49, 1373-1384.	9.4	783
48	[O1–11–03]: CEREBROSPINAL FLUID ENDOPHENOTYPES PROVIDE INSIGHT INTO BIOLOGY UNDERLYING ALZHEIMER'S DISEASE. Alzheimer's and Dementia, 2017, 13, P218.	0.4	0
49	Linkage, whole genome sequence, and biological data implicate variants in RAB10 in Alzheimer's disease resilience. Genome Medicine, 2017, 9, 100.	3.6	67
50	The Opioid Abuse Risk Screener predicts aberrant same-day urine drug tests and 1-year controlled substance database checks: A brief report. Health Psychology Open, 2017, 4, 205510291774845.	0.7	2
51	Fine-mapping of the human leukocyte antigen locus as a risk factor for Alzheimer disease: A case–control study. PLoS Medicine, 2017, 14, e1002272.	3.9	67
52	Discovery and Confirmation of Diagnostic Serum Lipid Biomarkers for Alzheimer's Disease Using Direct Infusion Mass Spectrometry. Journal of Alzheimer's Disease, 2017, 59, 277-290.	1.2	19
53	Seroprevalence and Serointensity of Latent Toxoplasma gondii in a Sample of Elderly Adults With and Without Alzheimer Disease. Alzheimer Disease and Associated Disorders, 2016, 30, 123-126.	0.6	33
54	Genetic studies of plasma analytes identify novel potential biomarkers for several complex traits. Scientific Reports, 2016 , 6 , $.$	1.6	25

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55	Genome-wide association study of prolactin levels in blood plasma and cerebrospinal fluid. BMC Genomics, 2016, 17, 436.	1.2	2
56	Assessment of the genetic variance of late-onset Alzheimer's disease. Neurobiology of Aging, 2016, 41, 200.e13-200.e20.	1.5	174
57	Crowdsourced estimation of cognitive decline and resilience in Alzheimer's disease. Alzheimer's and Dementia, 2016, 12, 645-653.	0.4	72
58	Shared genetic contribution to ischemic stroke and Alzheimer's disease. Annals of Neurology, 2016, 79, 739-747.	2.8	56
59	Presenilin E318G variant and Alzheimer's disease risk: the Cache County study. BMC Genomics, 2016, 17, 438.	1.2	11
60	Evaluating the necessity of PCR duplicate removal from next-generation sequencing data and a comparison of approaches. BMC Bioinformatics, 2016, 17, 239.	1.2	124
61	Variants in ACPP are associated with cerebrospinal fluid Prostatic Acid Phosphatase levels. BMC Genomics, 2016, 17, 439.	1.2	1
62	Variants in CCL16 are associated with blood plasma and cerebrospinal fluid CCL16 protein levels. BMC Genomics, 2016, 17, 437.	1.2	1
63	Genes for a â€`Wellderly' Life. Trends in Molecular Medicine, 2016, 22, 637-639.	3.5	7
64	A potential endophenotype for Alzheimer's disease: cerebrospinal fluid clusterin. Neurobiology of Aging, 2016, 37, 208.e1-208.e9.	1.5	44
65	Interaction between variants in <i>CLU</i> and <i>MS4A4E</i> modulates Alzheimer's disease risk. Alzheimer's and Dementia, 2016, 12, 121-129.	0.4	18
66	A novel Alzheimer disease locus located near the gene encoding tau protein. Molecular Psychiatry, 2016, 21, 108-117.	4.1	260
67	Influence of Coding Variability in APP-Aβ Metabolism Genes in Sporadic Alzheimer's Disease. PLoS ONE, 2016, 11, e0150079.	1.1	34
68	Discovery and Subsequent Confirmation of Novel Serum Biomarkers Diagnosing Alzheimer's Disease. Journal of Alzheimer's Disease, 2015, 49, 317-327.	1.2	10
69	Bridging the Gap between Statistical and Biological Epistasis in Alzheimer's Disease. BioMed Research International, 2015, 2015, 1-7.	0.9	26
70	Rarity of the Alzheimer Disease–Protective <i>APP</i> A673T Variant in the United States. JAMA Neurology, 2015, 72, 209.	4.5	41
71	Genetic studies of quantitative MCI and AD phenotypes in ADNI: Progress, opportunities, and plans. Alzheimer's and Dementia, 2015, 11, 792-814.	0.4	241
72	Genetically predicted body mass index and Alzheimer's disease–related phenotypes in three large samples: Mendelian randomization analyses. Alzheimer's and Dementia, 2015, 11, 1439-1451.	0.4	46

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73	Convergent genetic and expression data implicate immunity in Alzheimer's disease. Alzheimer's and Dementia, 2015, 11, 658-671.	0.4	173
74	Population-based analysis of cholesteryl ester transfer protein identifies association between I405V and cognitive decline: the Cache County Study. Neurobiology of Aging, 2015, 36, 547.e1-547.e3.	1.5	8
75	Genome-Wide Association Study of CSF Levels of 59 Alzheimer's Disease Candidate Proteins: Significant Associations with Proteins Involved in Amyloid Processing and Inflammation. PLoS Genetics, 2014, 10, e1004758.	1.5	109
76	A Versatile Omnibus Test for Detecting Mean and Variance Heterogeneity. Genetic Epidemiology, 2014, 38, 51-59.	0.6	52
77	Effects of Multiple Genetic Loci on Age at Onset in Late-Onset Alzheimer Disease. JAMA Neurology, 2014, 71, 1394.	4.5	166
78	Population-based Analysis of Alzheimer's Disease Risk Alleles Implicates Genetic Interactions. Biological Psychiatry, 2014, 75, 732-737.	0.7	52
79	Observed Changes in Radiographic Measurements of the First Ray after Frontal Plane Rotation of the First Metatarsal in a Cadaveric Foot Model. Journal of Foot and Ankle Surgery, 2014, 53, 274-278.	0.5	40
80	Genetic Discoveries in AD Using CSF Amyloid and Tau. Current Genetic Medicine Reports, 2014, 2, 23-29.	1.9	10
81	Genetic analysis of quantitative phenotypes in AD and MCI: imaging, cognition and biomarkers. Brain Imaging and Behavior, 2014, 8, 183-207.	1.1	161
82	Rare coding variants in the phospholipase D3 gene confer risk for Alzheimer's disease. Nature, 2014, 505, 550-554.	13.7	425
83	Variant Tool Chest: an improved tool to analyze and manipulate variant call format (VCF) files. BMC Bioinformatics, 2014, 15, S12.	1.2	11
84	Mitochondrial genomic variation associated with higher mitochondrial copy number: the Cache County Study on Memory Health and Aging. BMC Bioinformatics, 2014, 15, S6.	1.2	15
85	Population substructure in Cache County, Utah: the Cache County study. BMC Bioinformatics, 2014, 15, S8.	1.2	7
86	Observed Changes in First Metatarsal and Medial Cuneiform Positions after First Metatarsophalangeal Joint Arthrodesis. Journal of Foot and Ankle Surgery, 2014, 53, 32-35.	0.5	12
87	Phosphorylated Tau-AÎ ² 42 Ratio as a Continuous Trait for Biomarker Discovery for Early-Stage Alzheimer's Disease in Multiplex Immunoassay Panels of Cerebrospinal Fluid. Biological Psychiatry, 2014, 75, 723-731.	0.7	72
88	Missense variant in TREML2 protects against Alzheimer's disease. Neurobiology of Aging, 2014, 35, 1510.e19-1510.e26.	1.5	110
89	Variants in <i>PPP3R1</i> and <i>MAPT</i> are associated with more rapid functional decline in Alzheimer's disease: The Cache County Dementia Progression Study. Alzheimer's and Dementia, 2014, 10, 366-371.	0.4	36
90	Calibrating Longitudinal Cognition in Alzheimer's Disease Across Diverse Test Batteries and Datasets. Neuroepidemiology, 2014, 43, 194-205.	1.1	43

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91	Gene-Wide Analysis Detects Two New Susceptibility Genes for Alzheimer's Disease. PLoS ONE, 2014, 9, e94661.	1.1	155
92	Assessment of TREM2 rs75932628 association with Alzheimer's disease in a population-based sample: the Cache County Study. Neurobiology of Aging, 2013, 34, 2889.e11-2889.e13.	1.5	47
93	Meta-analysis of 74,046 individuals identifies 11 new susceptibility loci for Alzheimer's disease. Nature Genetics, 2013, 45, 1452-1458.	9.4	3,741
94	<i>TREM2</i> Variants in Alzheimer's Disease. New England Journal of Medicine, 2013, 368, 117-127.	13.9	2,385
95	O4-01-01: Association of genetic variants with cerebrospinal fluid protein levels of ACE, MMP3 and other proteins and risk for Alzheimer's disease., 2013, 9, P677-P678.		1
96	GWAS of Cerebrospinal Fluid Tau Levels Identifies Risk Variants for Alzheimer's Disease. Neuron, 2013, 78, 256-268.	3.8	344
97	Genetics of Alzheimer's Disease. BioMed Research International, 2013, 2013, 1-13.	0.9	75
98	Mitochondrial Haplotypes Associated with Biomarkers for Alzheimer's Disease. PLoS ONE, 2013, 8, e74158.	1.1	28
99	Alzheimer's Disease: Analyzing the Missing Heritability. PLoS ONE, 2013, 8, e79771.	1.1	257
100	Strong Evidence for a Genetic Contribution to Late-Onset Alzheimer's Disease Mortality: A Population-Based Study. PLoS ONE, 2013, 8, e77087.	1.1	14
101	The Role of Variation at \hat{A}^2 PP, PSEN1, PSEN2, and MAPT in Late Onset Alzheimer's Disease. Journal of Alzheimer's Disease, 2012, 28, 377-387.	1.2	53
102	Cerebrospinal fluid APOE levels: an endophenotype for genetic studies for Alzheimer's disease. Human Molecular Genetics, 2012, 21, 4558-4571.	1.4	196
103	Novel late-onset Alzheimer disease loci variants associate with brain gene expression. Neurology, 2012, 79, 221-228.	1.5	144
104	Single nucleotide polymorphism discovery in cutthroat trout subspecies using genome reduction, barcoding, and 454 pyro-sequencing. BMC Genomics, 2012, 13, 724.	1.2	18
105	Mitochondrial Genomic Analysis of Late Onset Alzheimer's Disease Reveals Protective Haplogroups H6A1A/H6A1B: The Cache County Study on Memory in Aging. PLoS ONE, 2012, 7, e45134.	1.1	44
106	Fine Mapping of Genetic Variants in BIN1, CLU, CR1 and PICALM for Association with Cerebrospinal Fluid Biomarkers for Alzheimer's Disease. PLoS ONE, 2011, 6, e15918.	1.1	64
107	Common variants at MS4A4/MS4A6E, CD2AP, CD33 and EPHA1 are associated with late-onset Alzheimer's disease. Nature Genetics, 2011, 43, 436-441.	9.4	1,676
108	Common variants at ABCA7, MS4A6A/MS4A4E, EPHA1, CD33 and CD2AP are associated with Alzheimer's disease. Nature Genetics, 2011, 43, 429-435.	9.4	1,708

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109	Performances on the CogState and Standard Neuropsychological Batteries Among HIV Patients Without Dementia. AIDS and Behavior, 2011, 15, 1902-1909.	1.4	52
110	Association and Expression Analyses With Single-Nucleotide Polymorphisms in <emph type="ital">TOMM40</emph> in Alzheimer Disease. Archives of Neurology, 2011, 68, 1013.	4.9	97
111	SNPs Associated with Cerebrospinal Fluid Phospho-Tau Levels Influence Rate of Decline in Alzheimer's Disease. PLoS Genetics, 2010, 6, e1001101.	1.5	111
112	Genetic Evidence Implicates the Immune System and Cholesterol Metabolism in the Aetiology of Alzheimer's Disease. PLoS ONE, 2010, 5, e13950.	1.1	347
113	Validating predicted biological effects of Alzheimer's disease associated SNPs using CSF biomarker levels. Journal of Alzheimer's Disease, 2010, 21, 833-42.	1.2	43
114	Rs5848 Variant Influences GRN mRNA Levels in Brain and Peripheral Mononuclear Cells in Patients with Alzheimer's Disease. Journal of Alzheimer's Disease, 2009, 18, 603-612.	1.2	59
115	Alzheimer's disease risk variants show association with cerebrospinal fluid amyloid beta. Neurogenetics, 2009, 10, 13-17.	0.7	80
116	Genome-wide association study identifies variants at CLU and PICALM associated with Alzheimer's disease. Nature Genetics, 2009, 41, 1088-1093.	9.4	2,697
117	Molecular characterization of novel progranulin (<i>GRN</i>) mutations in frontotemporal dementia. Human Mutation, 2008, 29, 512-521.	1.1	71
118	Novel presenilin 1 variant (P117A) causing Alzheimer's disease in the fourth decade of life. Neuroscience Letters, 2008, 438, 257-259.	1.0	14
119	Variation in <i>MAPT</i> is associated with cerebrospinal fluid tau levels in the presence of amyloid-beta deposition. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 8050-8054.	3.3	84
120	Association studies between common variants in prolyl isomerase Pin1 and the risk for late-onset Alzheimer's disease. Neuroscience Letters, 2007, 419, 15-17.	1.0	25
121	Association studies testing for risk for late-onset Alzheimer's disease with common variants in the \hat{l}^2 -amyloid precursor protein (APP). American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 469-474.	1.1	11
122	Extreme cerebrospinal fluid amyloid \hat{l}^2 levels identify family with late-onset Alzheimer's disease presenilin 1 mutation. Annals of Neurology, 2007, 61, 446-453.	2.8	87
123	Haplotype-based association analysis of the MAPT locus in late onset Alzheimer's disease. BMC Genetics, 2007, 8, 3.	2.7	45
124	Apolipoprotein E levels in cerebrospinal fluid and the effects of ABCA1 polymorphisms. Molecular Neurodegeneration, 2007, 2, 7.	4.4	68
125	Identification and validation of novel CSF biomarkers for early stages of Alzheimer's disease. Proteomics - Clinical Applications, 2007, 1, 1373-1384.	0.8	66
126	A Scan of Chromosome 10 Identifies a Novel Locus Showing Strong Association with Late-Onset Alzheimer Disease. American Journal of Human Genetics, 2006, 78, 78-88.	2.6	157

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127	Ubiquilin 1 polymorphisms are not associated with late-onset Alzheimer's disease. Annals of Neurology, 2006, 59, 21-26.	2.8	37
128	HDDD2 is a familial frontotemporal lobar degeneration with ubiquitin-positive, tau-negative inclusions caused by a missense mutation in the signal peptide of progranulin. Annals of Neurology, 2006, 60, 314-322.	2.8	186
129	DAPK1 variants are associated with Alzheimer's disease and allele-specific expression. Human Molecular Genetics, 2006, 15, 2560-2568.	1.4	125
130	Association studies between risk for late-onset Alzheimer's disease and variants in insulin degrading enzyme. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2005, 136B, 62-68.	1.1	35
131	Microsatellites versus single-nucleotide polymorphisms in linkage analysis for quantitative and qualitative measures. BMC Genetics, 2005, 6, S122.	2.7	15
132	An analysis of identical single-nucleotide polymorphisms genotyped by two different platforms. BMC Genetics, 2005, 6, S152.	2.7	6
133	The efficacy of short tandem repeat polymorphisms versus single-nucleotide polymorphisms for resolving population structure. BMC Genetics, 2005, 6, S84.	2.7	7
134	Association of late-onset Alzheimer's disease with genetic variation in multiple members of the GAPD gene family. Proceedings of the National Academy of Sciences of the United States of America, 2004, 101, 15688-15693.	3.3	134
135	Phylogeographic and nested clade analysis of the stonefly Pteronarcys californica (Plecoptera:Pteronarcyidae) in the western USA. Journal of the North American Benthological Society, 2004, 23, 824-838.	3.0	27
136	Genome assembly of the roundjaw bonefish (Albula glossodonta), aÂvulnerable circumtropical sportfish. GigaByte, 0, 2022, 1-29.	0.0	1